

# 1030 at Structural Genomics

- lighting is largest challenge
  - saturation of top or bottom lighting
  - lensing at top drop and liquid in well
- Drop is 50% test liquid and 50% liquid
  - well has liquid and standard thermodynamics
  - liquid migrates to well out of drop
  - leave solution behind with hopefully crystals.

*Optical  
machines*

- Emerald biosciences
  - Hardware + software
- Cytoe labs → Gibson → Chunky hardware is slow
  - Hardware + SW out of U of Alabama

0	Clear
1	light
2	Medium / Heavy
3	Ugly
4	Phase separation
5	?
6	Spherulites
7	Grainy precipitate
8	Micr crystals < 20 μm
9	Crystals > 20 μm

to catalog via video

A1 → 0

A2 → 3

A3 → 7

" "

" "

" "

lighting JR filter

back light

top light

Autofocus lens or manual 2mm thick drop size

Scanned plates upper right corner checker

Auto zoom - focus w/ focus, light change at zooms  
 3x, 10x, 20x

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Enclosure 3